

## CLAIMS

1. A method of infecting the glomerular cells of a kidney of a mammalian subject  
5 requiring same with a recombinant adenovirus vector carrying a gene or genes of interest,  
comprising the step of infusing intra-renal arterially in a single pass through the superior  
mesenteric artery or renal artery an effective amount of said adenoviral vector into said  
kidney at an effectively slow rate over an effective period of time, under conditions such  
that at least 30% of said glomerular cells are infected with said vector.

2. The method according to claim 1, wherein said adenovirus vector carries a control  
element that preferentially expresses said gene or genes into renal glomerular cells.

3. The method according to claim 1, wherein said kidney is maintained at reduced  
temperatures during said infusion procedure,

4. The method according to claim 1, further comprising clamping the aorta above and  
below said superior mesenteric renal artery of said kidney, and infusing through said  
superior mesenteric renal artery.

5. The method of claim 1, wherein said renal artery is cannulated directly without  
clamping of said aorta during said infusion.

6. The method of claim 1, wherein said mammal is a rodent, said rate of infusion is  
about  $0.1-0.5 \times 10^{11}$  particles per minute, and said effective period of adenoviral vector  
infusion is between about 15 and 120 minutes.

7. The method of any one of claims 1 through 6, further comprising concurrent  
cannulation of the femoral vein through the vena cava into the renal vein so as to direct  
vector not taken up by renal glomerular cells away from the general circulation.